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ANTIMICROBIAL PROTEINS FROM THE SPO1 BACTERIOPHAGE

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ANTIMICROBIAL PROTEINS FROM THE SPO1 BACTERIOPHAGE

CROSS-REFERENCE TO RELATED APPLICATIONS

[01] This Application claims priority to provisional application 60/457287 filed on March 25, 2003.

STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH OR DEVELOPMENT

[02] The present invention may have been developed with funds from the United States Government. Therefore, the United States Government may have certain rights in the invention.

REFERENCE TO A COMPACT DISK APPENDIX

[03] A Sequence Listing, including SEQ ID NO: 1 to 24, is submitted with this application.

FIELD OF THE INVENTION

[04] The invention relates to proteins from the SPO1 bacteriophage, genes encoding same, and their use in arresting bacterial metabolism. Further, the present invention relates to a method of treating infected patients through the introduction of proteins, peptide fragments of the proteins or peptidomimetics, or genes encoding same into the bacteria infecting the patient.

BACKGROUND OF THE INVENTION

[05] Viruses are obligate parasites that use the host's mechanisms for expressing their genes and propagating themselves. Bacteriophage, or "phage" for short, are viruses that infect bacteria. Phage have a host range that is often fairly narrow, infecting a single species or a group of related species.

[06] Some bacteriophage, such as T4, always kill their hosts within a short time after infection--they are called LYTIC phages. Others, such as Lambda, can also establish themselves in a dormant form within the cell and be maintained stably over many generations--they are called LYSOGENIC phages. On infection lysogenic phages have a choice between a lytic cycle or to establish lysogeny.

[07] In a lysogenic cycle, the phage integrates into the host chromosome at a specific point using a site-specific recombination process. The expression of most phage genes is shut down by a repressor and the phage genome can be replicated as part of the host chromosome. Under certain conditions, usually involving host stress, the genes get switched on again and the phage genome is excised from the chromosome, replicated and new phage particles are released by lysing the cell.

[08] After their discovery early in the 20th century, phage were widely used to treat various bacterial diseases in people and animals. After an enthusiastic beginning, poorly designed research protocols and the development of antibiotics ended most phage therapy research in the West. However, phage therapy continued in Poland and the former Soviet Union for decades and the Eastern European experience should serve to bootstrap the technology into the U.S. Thus, although there are few, if any, clinical trials underway in the U.S., the technology is reasonably well developed and its advantages and limitations are well understood.

[09] The alarming rise in antibiotic resistance in bacteria is leading to a resurgence of interest in bacteriophage research. In fact several companies exist that are dedicated to the realization of phage therapy. The Eliava Institute of Bacteriophage, Microbiology & Virology in the Republic of Georgia has been an international leader in phage therapy for over 70 years and has the world's largest collection of therapeutic phages. Other companies including Exponential Biotherapies, Inc. of New York; Intralytix, Inc. in Maryland; Phage Therapeutics International, Inc. (PhageTx) of British Columbia; Biophage, Inc. of Quebec; and PhageTech, Inc. of Quebec are developing commercial phage therapies.

[10] The limitations and advantages of phage therapy are listed by Elizabeth Kutter of Evergreen State College at <http://www.evergreen.edu/phage/phagetherapy.html>. Her lists are worth iterating here in order to understand the potential and limitations of phage therapy. The limitations

include: 1) Paucity of understanding of the heterogeneity and ecology of both the phages and the bacteria involved; 2) Failure to select phages of high virulence against the target bacteria before using them in patients; 3) Use of single phages in infections that involve mixtures of different bacteria; 4) Emergence of resistant bacterial strains, which can occur by selection of resistant mutants (a frequent occurrence if only one phage strain is used against a particular bacterium) or by lysogenization (if temperate phages are used); 5) Failure to appropriately characterize or titer phage preparations, some of which were totally inactive; 6) Failure to neutralize gastric pH prior to oral phage administration; 7) Inactivation of phages by both specific and nonspecific factors in body fluids; 8) Liberation of endotoxins as a consequence of widespread lysis of bacteria in the body (this is called the Herxheimer reaction); and 9) Failure to identify the bacterial pathogens involved necessitated by the relative specificity of phage therapy.

[11] The advantages of phage therapy include: 1) They are self-replicating and self-limiting, because they multiply only as long as sensitive bacteria are present and then are gradually eliminated; 2) They target specific bacteria, causing less damage to the normal microbial balance in the body; 3) Phage can often be targeted to receptors on the bacterial surface that are involved in pathogenesis, so that any resistant mutants are attenuated in virulence; 4) Few side effects have been reported for phage therapy; 5) Phage therapy would be particularly useful for people with allergies to antibiotics; 6) Appropriately selected phages can easily be used to prevent bacterial disease in people or animals at times of exposure, or to sanitize hospitals and help protect against hospital-acquired infections; 7) Phage can be prepared fairly inexpensively and locally, facilitating their potential applications to underserved populations; 8) Phage can be used independently or in conjunction with other antibiotics to help reduce the development of bacterial resistance; and 9) Multiple delivery means are available, including noninvasive means like topical application, oral administration, and inhalation.

[12] One lytic phage of long time interest to our laboratory is called "SPO1." SPO1 has a linear dsDNA genome of 140 kb, and its host is the bacterium *Bacillus subtilis*. Early in SPO1 infection of *B. subtilis*, the synthesis of most host-specific molecules is replaced by the corresponding phage-specific biosyntheses. Subversion of the host machinery is accomplished primarily by a cluster of early genes in the SPO1 terminal repeat in an 11.5-kb "host-takeover module." The module includes

24 genes, tightly packed into 12 operons driven by the previously identified early promoters PE1 to PE12.

[13] The 24 genes are smaller than average, with half of them having fewer than 100 codons. Most of their inferred products show little similarity to known proteins, although zinc finger, trans-membrane, and RNA polymerase-binding domains were identified therein. Transcription-termination and RNase III cleavage sites were identified in the nucleotide sequence as well. We have placed most of these 24 genes into an inducible *B. subtilis*/*E. coli* shuttle vector and introduce them here for use in phage therapy and as antimicrobial agents.

BRIEF SUMMARY OF THE INVENTION

[14] As used herein "tightly regulated inducible promoter" is an inducible promoter that does not express detectable levels of protein in the absence of the inducing agent. Detectable levels of protein expression could be indicated by inhibition of bacterial host cell growth. In other words, the promoter is not "leaky" to a degree sufficient to prevent cloning of the gene in the host cell.

[15] As used herein "recombinant" is relating to, derived from, or containing genetically engineered material.

[16] "% identity" is calculated over the entire length of a specified sequence, thus short local alignments with a claimed sequence are not relevant (e.g., % identity = number of aligned residues/length of reference sequence). Alignments are performed using BLAST (Basic Local Alignment Search Tool) homology alignment as described by Tatusova and Madden (1999). The default parameters are used, except the filters are turned OFF. As of Jan. 1, 2001 the default parameters were as follows: BLASTN or BLASTP as appropriate; Matrix = none for BLASTN, BLOSUM62 for BLASTP; G Cost to open gap default = 5 for nucleotides, 11 for proteins; E Cost to extend gap [Integer] default = 2 for nucleotides, 1 for proteins; q Penalty for nucleotide mismatch [Integer] default = -3; r reward for nucleotide match [Integer] default = 1; e expect value [Real] default = 10; W wordsize [Integer] default = 11 for nucleotides, 3 for proteins; y Dropoff (X) for blast extensions in bits (default if zero) default = 20 for BLASTN, 7 for other programs; X dropoff value for gapped alignment (in bits) 30 for BLASTN, 15 for other programs; Z final X dropoff value for gapped alignment (in bits) 50 for

BLASTN, 25 for other programs. This program is available online at <http://www.ncbi.nlm.nih.gov/BLAST/>.

[17] The invention provides expression vectors encoding a protein comprising a sequence selected from the group consisting of SEQ ID NO: 1-24, wherein the expression of the protein is controlled by an inducible promoter. The transcription must be under sufficiently tight control so as to allow the vector to be propagated in the host cell.

[18] The invention also provides a method of designing peptidomimetic anti-bacterial drugs comprising preparing proteins of SEQ ID NO: 1 to 24 or variants with at least 95% identity over the entire length of the specified sequence by recombinant or synthetic means (e.g., peptide synthesis), determining their three-dimensional structures, and using the three-dimensional structure as the basis for designing peptidomimetic small molecules to be tested for anti-bacterial activity.

[19] Methods of treating a bacterial infection in a warm blooded animal in need thereof are also provided. The methods require the direct *in vivo* delivery of an antimicrobial protein as described above, in an amount effective to kill bacteria, or delivery of a peptidomimetic small molecule having the bactericidal activity of the protein. Alternatively, the protein can be indirectly delivered by a bacteriophage that encodes said protein. Beneficially, the proteins of the invention do not lyse the bacteria, and are of particular use where an infection is halted without cell lysis and the concomitant release of bacterial toxins.

BRIEF DESCRIPTION OF THE DRAWINGS

[20] FIG. 1. Shutoff of Host RNA Synthesis. At various times after infection at 30°C, cultures were pulse-labeled with [5-³H] uridine. Single mutants 50- and 51- had no significant effect (data not shown), showing that both gene products, GP50 and GP51, contribute to the activity of GP50/51 ("GP" refers to "gene product" throughout). Each data point in such an experiment represents the sum of host and phage RNA synthesis.

[21] FIG. 2. Shutoff of Host DNA Synthesis. At various times after infection at 30°C, cultures were pulse-labeled with [methyl-³H] thymidine. Only host DNA synthesis was assayed in such an experiment, since SPO1 incorporates hmUra into its DNA in place of thymine.

[22] FIG. 3. Shutoff of Host RNA Synthesis. Host and phage RNA synthesis were monitored as described for 39-, 40-, and 39-40- SPO1.

[23] FIG. 4. Shutoff of Host DNA Synthesis. Host DNA synthesis was monitored in 39-, 40-, and 39-40- SPO1.

[24] FIG. 5. Gene-Specific Expression Patterns. Cultures infected with wild-type SPO1, mutant 28-, or triple mutant 44-50-51- were harvested at 5, 15, or 25 minutes after infection at 30°C. Triangles represent wild-type; squares represent the 28- mutant; and circles the triple mutant 44-50-51-.

[25] FIG. 6. Differential Effects of Mutations on Immediate-Early, Delayed-Early, and Middle Transcription. Cultures were infected with wild-type SPO1, mutant 44-, double mutant 50-51-, or triple mutant 44-50-51-. The numbers at the top represent the gene or genes present in that spot on the dot blot. The gene categories are indicated across the top. Gene 27 falls under both middle and delayed-early categories.

[26] FIG. 7A. Distinguishable Effects of Genes 50 and 51 in the presence of the 44- mutation. Rates of RNA synthesis were measured by pulse-labeling with [5-³H]uridine.

[27] FIG. 7B. Distinguishable Effects of Genes 50 and 51 in the absence of the 44- mutation. Rates of RNA synthesis were measured by pulse-labeling with [5-³H]uridine.

[28] FIG. 8. Expression of SPO1 GP44 in *E. coli*. Cells of *E. coli* strain DH5(pPW27), which expresses SPO1 gene 44 from an IPTG-inducible promoter, were transformed either with vector pACYC177 or with plasmid pPW40, in which the mutant *rpoB* gene had been cloned in pACYC177. Aliquots of the two cultures were induced by adding 0.2 mM IPTG at time 0, and were assayed for viable cells at various times thereafter, by plating on media lacking IPTG. Open symbols, no IPTG; Closed symbols, 0.2 mM IPTG added at time 0; Squares, mutant *rpoB*; and Circles, no mutant *rpoB*. Viable cell count is expressed in cells/ml X 10⁻⁶.

[29] FIG. 9. Effect of Gene 40, Gene 44, Gene 50 and Gene 51 on Phage DNA Synthesis.

[30] FIG. 10. Effect of Gene 44, Gene 50 and Gene 51 on Single Step Growth Curves.

[31] FIG. 11. Effect of Gene 39 and Gene 40 on Single Step Growth Curves.

[32] FIG. 12. Effect of GP50 on Host DNA Synthesis.

DETAILED DESCRIPTION OF THE INVENTION

[33] The early gene region of the terminal redundancy of SPO1 is found at GenBank Acc. No. AF031901. It is a linear 11500 bp dsDNA, where 5-hydroxymethyluracil (hmUra) is used instead of thymine as the base-pairing partner for adenine, and T's in the GenBank sequence represent hmUra. There are 24 small proteins encoded therein. The 24 genes are arranged in 12 operons, each of which includes between one and 4 genes. Genes 51 and 50 constitute one operon; gene 44 is the only gene in its operon; genes 37-40 are all in one operon. Genes 37-49 are transcribed rightward and 60-50 are transcribed leftward, which is why the latter are expressed in reverse numerical order. The genes and their encoded proteins are listed in Table 1.

Table 1 - Anti-bacterial proteins from SPO1		
Gene Name	SEQ ID NO:	Peptide Sequence
37	1	MSDVIIIPFLTSAVTAFIVAYLLDRWYIKRRR
38	2	MSVQIKHGNKTFVVDPSGDVKEGSYVLYLYEYRLGEVDVGRVSEVANDGRLYLDGPGVIVTLDQ PFILLKEVVEEEDEDDRIDAEFHNPLLRKLENTTEKLTPEETQLAQWTTMTRVFSHDLKKGI PYAIKHKNSGNILYGLYSGLLNPTALFRHLNEESKISIEQLKSGLEIYEVVEDEEESIWN
39	3	MELNLDIYVDYKDKRYKAEGYYGPSVGDVLIFMDMELEGATVQEVARIEGSEIHLRTPNGNEP SYRYMGQYLILKPYGSSDPRGDILVHEDVQYVRVDAQAMPGLIEALEPNKLPFSGKRPFKYRPA VLEVEYVLTKDEQVLQLENGKSYSGAYRVLI PRMGVLPKTHIYTHKHVFMEDVFLGNSYEL SSPNDVEMTPIHAVFTGFSGKNRDEAIFVNPYYNDGVTGTMITVSDLLTGKWDITPLVPKKGV
40	4	MHIYTYWGLKYVPSNSTMVAKEGDLILLGNEVHKVVKVLHRFRNITDLQITNWKGTETRYNLHV TEYKVLVPYDTHKEENEAMSDSLITHNGKDYVLCKIPARVGD LIRTEDKRVWEVLQSKDGLVL YNEEKGEQRSAYVSEIGPYHVLVPRDTHPTREELAAVIMNKAFTRTETQDSQEDTGTHKGL GLTGTDLYHSLRDLDAKVQSGYYTATENEEDVKSEIEATKKHMKAVKESGKTVNDYRKEENTKR CKLKALTNKFNRLFLKSVIDTDSLQVGKAYLIGGRDMKNVHGLYTGTTFDQQHANFLIVETDRM HRTLTVSAEQLF AEERHIVDIEKRVEQTED
41	5	MEKLPNTVVKVHGEEMESKLPFRKLHKDTNSILREDLVSACQEHIEALVEGMIAHGDGRKVAEL DTSTQYYWHLKLVEYTPIPGRQTQHYVDLVDGTNPVCYFSLCDCSGDNI TDRRWSNMVKRLQNP EEDIAKTLRCYFRQDAGMPSWIEYPQ
42	6	MRKFVTTLTASPRNKKVGNHRLEISPFVSLRRYYYFNTAICIENPVTREFAIDDSYGSLSTNQN CAQYRQYFSLGGYKEVSLEEIHAV
43	7	MIQLSERQQDLLQVAEKYEQCHIEFYTAQSRLFGTEIMGEVVKTSGLTKIAHPEEDLFEVALA YLASKKDILTAQERKDVLFYIQNNLC
44	8	MAKSNNVYVVGEEKVSTLAEVAKVLGVSRSVKDVEEGKYDVVVEEAAVSLADTEEVVEEVVT

Table 1 - Anti-bacterial proteins from SPO1		
Gene Name	SEQ ID NO:	Peptide Sequence
		EEEDILEGVEVVEDEEEEEAAEDVEEPTSEEDSEDEWEEGYPVATEVEEDEDEEIEYPEVGDFE DEKAIKKYIKGLTDEQLQAWCELEGAEWVENEHRNINRMRMAMAIAVHFPELAKKPSSKKKSK YA EYTTEELVEMAIDNNVEVRDDKGNERILRMYTIIALREAGLIS
45	9	MMMDKQVEEVKKHYPIVEDWSVIVARKEDDCMTVTDAVPFILAGYKNVSYEMDDIVVLCSEPIG LTWEDVRFLKNHEGSVSFEEIGYEDKAMVYHVDLG
46	10	MMTEDQKFYLTKEIELEAGCFSDWTKEDITGDLKYLKKGII EESIELIRAVNGLTYSEELHDF TQEII EELDISPL
47	11	MDWTKMTFMGTVDVEKWIWNGLEEAGRLYAVWLSDDHVGIVDVNEEGLFCLGWVSDISPESLQ NMLGGGAELFESYEDVLSEHGGSIAIRVEV
48	12	MPYSKITVPVLVGEGLTEWDVIDVMRETHPPTVEDQYHYHTFDSMQNRTIFVLENPLYPDVDKI PEKVLGIAVDALEDMLDNVPVEDLPVTEEQGNVKRFTTKLASIVFDVFLIIPDFVSVTAKEE
49	13	MIKAAVTKESLYRMNTLMEAFQGFGLGLDGEFTFKVKPGVFLITDVKSYLEGDKYDDAFNALID FVLRNDRDAVEGTETDVSIRLGLSPSDMVVKRQDKTFTFTHGDLFEVHWINL
50	14	MDKLAAGGLYLLFLLLAGIIVTH
51	15	MAKINKGYVANFIEENGFPQQGHFEKKDLQAFYKHLSTEQLEEWVELEGLEVKDTSDSDSIYRM RLCMAILYLNFPKKTAGKKKASPYKHISLEELVQMATDNDIEVKHTDSDKILRMRTIMALKEAG KLG
52	16	MTHFISIATYIYALVSAGFIGGWHEESWIKDTEYEHGGYHMIIDTPAVVNYSLEYGNYQWIFQ KYMKEGKVTVERFYRNSLDIPKEILTDEALAFIKWDENANEYELHAGEGVLYFKYEGEEKGYV IPMAYAGEIMFVPDEDAEKALEIINSQKKY
53	17	MRTYWNVSLDRSNGKRFRERLVHYICVPIISIHHAEDTISMTRKEVGH LAETIANHIILDINGTY RTFSVNDIVHCSLEKVITLEGDVTNEFIDRLQILVNKEVQGSQSTQQSLSSVFESTLEKYN SPD DFADYLEETEEEDYEDYSLDDTIDAISYALKTQEPVQAEWCCLMVDVYTGTLTEVTVETDKDK TLDSILGKYLENGFECVSKRLGEVL
54	18	MVIIKYTTKTQPTPVKEMFISPQHYAKWRSHMGSKLTSVKPIKGR
55	19	MFKLLTLFKRNKITSAAEYYTQAIHICEQFDRSTQKYTSM
56	20	MFKYTDRSVRQYIERQORSAMLEQEQAEDKKERRKAGLLFFGTIVVLVAVVAVYIVPQSLDAM WHENYEKPAQEAARN
57	21	MTLFIAGVTLEEVREATVSALFVKLEQEKALYLGAGSEDSNLCKSTLDKVQEDYPLDDMEKD YLRDLLQFWLSRLFLGDGFEGEIPDSSEDLRRTATTAFTYTAAIRHYCM
58	22	MTLAGYRVDSCNGCGKAYLVGESHDRKKCAECASK
59	23	MKKRYKVTFEDGTSQCLVVGNFSSPTNAWCAAMRNLTPEGIARVQHYNVEEISK
60	24	LNQVEVLREEYVEGYVVMWRNPSNAPVIEVFTEDNLEEGIIPEYVTANDDTFDRIVDAVEFG YLEEELV

[34] **Single Step Growth Experiment:** Cells of CB10 were infected at an MOI of about 0.1 at 30°C. The infected cultures were diluted through anti-SPO1 antibody and plated on CB313 lawns at 25 minute intervals. Plaque forming units (Pfu) were counted and plotted against time.

[35] **SDS-PAGE:** Cultures of CB10 were infected with appropriate strains of SPO1 at an MOI of 5, and shaken at 30°C. At various times, aliquots were pulse labeled for 2 minutes with (³⁵S) Met-Cys. Extracted proteins were subjected to SDS-PAGE, and autoradiographs were prepared.

[36] **RNA preparation:** Cultures in VY medium were harvested at the indicated times after infection, as described previously (Wei and Stewart, 1993). The RNA was purified using the QIAGEN® RNEASY® Mini Kit, following the manufacturer's protocol.

[37] **Dot blots:** PCR products carrying a particular gene or genes were denatured by incubating for 15 minutes at 37°C in 0.2M NaOH. 10 ng of DNA in 2 µl were spotted onto positively charged nylon membranes, AMERSHAM® HYBOND™ N+, and bound by UV crosslinking. Each 11 cm X 5 cm membrane held an array of about 23 spots. The PCR fragments included all, or nearly all, of the specified gene or genes, except for gene 31 which included only the first exon.

[38] **Hybridization:** For each RNA preparation, a 1.0 µg aliquot was labeled with alkaline phosphatase, using the AMERSHAM® ALKPHOS™ Direct Labeling and Detection Kit. The entire labeled preparation was hybridized to one membrane carrying a dot blot array, using the hybridization buffer defined in the kit. Hybridization was at 60°C for two hours in a ROBBINS® Model 400 hybridization incubator rotating at about 4 rpm. Wash buffers were as defined in the kit, with the primary wash at 65°C and the secondary wash at room temperature. Detection of hybridized RNA used the AMERSHAM® CDP-STAR™ Chemiluminescent Detection Reagent, following procedures from the kit. The alkaline phosphatase catalyses decomposition of dioxetane with emission of light, which is detected on film, using HYPERFILM ECL™ from AMERSHAM®. All hybridizations were done under the same conditions, without attempting to identify optimal conditions for each gene, and we have not made independent measurements of the concentrations of the various transcripts or of their intrinsic hybridization efficiencies. Thus, gene-to-gene differences in the absolute amount of signal detected could be due to differences in intrinsic hybridization or labeling efficiency. Conclusions were drawn only from comparisons between signal intensities for the same gene under different conditions.

[39] **Densitometry:** The intensities of the dot-blot signals were measured with a MOLECULAR DYNAMICS™ computing densitometer. Positions at which no spot was visible on the film gave intensity values ranging from -20 to +20. Some spots that were faintly visible gave intensity readings less than 20, which are reported as measured, with the recognition that they represent only approximations of the actual value. Spots not visible on film are reported as 0. Intensity values as high as 1700 were within the linear dose-response range. Where undiluted samples gave values beyond the

linear range, we measured hybridization by diluted samples, and corrected for the dilution factor and the experiment-to-experiment difference in hybridization efficiency

[40] **Pulse Labeling:** Rates of synthesis of host and phage RNA or protein were measured in C4 medium as described previously (Wei and Stewart, 1993), by pulse-labeling for 5 minutes with [5-³H]uridine, or [4,5-³H]leucine, precipitating with TCA, and counting the precipitate. Since SPO1 DNA contains hmUra in place of thymine, host DNA synthesis could be measured independently by pulsing with [methyl-³H]thymidine. Phage DNA synthesis was measured by pulsing with [2,8-³H]adenine, and measuring alkali-stable, TCA-precipitable counts. Pulsing with [2,8-³H]adenine measures host as well as phage DNA synthesis, but most host DNA synthesis has been shut off by the time that phage replication begins, even in mutant infections, so incorporation of label after 12 minutes is overwhelmingly into phage DNA. The time given for each pulse is the time at which the pulse began.

[41] **Mutagenesis:** Cloned genes were mutated using the STRATAGENE® QUIKCHANGE® kit. In each case, a nonsense codon was substituted for the first or second lysine codon (or for the first leucine codon in the case of gene 37) (codons 9, 10, 14, 10, 3, 11, 7, 3, 3, and 3 for genes 37, 38, 39, 40, 44, 45, 46, 50, 51 and 56, respectively), and the mutants were propagated on a suppressor strain (CB313) which inserts lysine at nonsense codons. Each mutagenized plasmid was allowed to recombine with superinfecting SPO1. Recombinant progeny were identified by plaque-lift hybridization, using the AMERSHAM ECL™ 3'-oligo labeling and detection kit, and allowed to segregate pure mutant strains. Multiple mutants were produced by successive recombination with the appropriate mutagenized plasmids and all genotypes were confirmed by DNA sequencing. The first 8 nucleotides of gene 50 overlap the last 8 nucleotides of gene 51, and the gene 50 mutation destroys the termination codon of gene 51, apparently permitting translation of gene 51 to run on for about 30 more amino acids, beyond its normal 151. Thus, definitive conclusions about the role of GP50 could not be made from single mutants in which only gene 50 was mutated. The gene 28 mutant was derived from mutant F21, initially described by Okubo *et al.* (1972), by 5 successive back crosses to wild-type SPO1. We have previously called it sus28-1 and our lab bookkeeping lists it as F21(5).

[42] Identification of Immediate-Early, Delayed-Early, and Middle Genes:

Expression of specific genes was measured by dot blot hybridization and densitometry. FIG. 5 shows the expression pattern of the known early genes and several representative middle genes, in the wild-type or when altered by either the 28- mutation or the 44-50-51- triple mutant. The 28- mutation inactivates the middle gene-specific sigma, permitting identification of middle gene activity. Note that different genes are expressed on different scales, as the absolute level of expression varies widely from gene to gene. The densitometric values for each gene are plotted as a function of time after infection. In each graph, the number in the upper left corner indicates the gene whose activity is plotted. Where an operon of more than one gene was assayed, the graph is labeled with the number of the first gene in the operon. This applies to operons 37-39, 45-46, 48-49, 51-50, 55-53, 58-56, and 60-59. To a first approximation, the immediate-early genes have been arranged from top to bottom in order of increasing activity. Gene 27 could have been displayed with either the delayed-early or the middle genes.

[43] Genes 31 and TF1 were previously known to be middle genes (Greene *et al.*, 1984; Scarlato and Gargano, 1992). The others are now identified as immediate-early genes (41-51 and 53-60), delayed-early genes (27, 28, and 37-40), or middle gene (52), on the basis of expression time in wild-type infection and of the GP28-dependence of their expression. Gene 27 had characteristics of both delayed-early and middle genes. Genes 48-49 also showed substantial GP28-dependent expression at late times (data not shown).

[44] For FIG 6, cells were harvested 15 minutes after infection at 30°C. RNA was prepared, labeled, hybridized to dot blots, and detected on film. The films were scanned using the UMAX® VISTA SCAN™ program with a UMAX® ASTRA 1220S™ scanner. A 25 X 25 pixel square surrounding each spot was copied and pasted into the grid displayed.

EXAMPLE 1: ANTIMICROBIAL ACTIVITY

[45] For analysis of antimicrobial activity we first cloned most of the 24 host-takeover genes (genes 37-60) singly or in multiple gene combinations into a *B. subtilis*/*E. coli* shuttle vector (pPW19) under control of an IPTG-inducible promoter (Wei and Stewart, 1993). Certain genes and gene combinations were not clonable using this vector system, suggesting that these genes are highly toxic, and that the small amount of leaky expression that occurs, even without induction, is sufficient to

kill the bacteria. These remaining genes are being cloned using expression vectors with more tightly regulated promoters, such as pJONEX4 (Sayers, 1996), pET101, or pX (Kim *et al.*, 1996). The transformed *E. coli* and *B. subtilis* were used to determine the antimicrobial activity, if any, of the cloned genes. The effect of induction on bacterial growth and viability was measured both in liquid medium and by colony formation. Definitive establishment of lethality was shown by loss of the ability to form colonies even after removal from the inducing agent. Indication of probable lethality was based on total inability to form colonies under inducing conditions. Data obtained to date are summarized in Table 2, and discussed in more detail below.

Table 2. Summary of Lethality Data					
Gene	<i>E. coli</i>	<i>B. subtilis</i>	Gene	<i>E. coli</i>	<i>B. subtilis</i>
37	Lethal *	No data	46	Lethal	Mildly Toxic
38	Highly toxic	No effect **	45/46	Highly Toxic	Lethal
39	Lethal	No data	47	No effect	No effect
38/39	Lethal	Lethal *	48	Strongly Toxic	Strongly Toxic
40	Mildly Toxic	Inconsistent	49	No data	No data
38/39/40	Lethal *	No data	48/49	Non-Toxic **	No data
41	Mildly Toxic	Lethal *	52	Lethal	Not tested
42	No effect	No effect	53/54/55	Toxic	Toxic
43	Indeterminate	Indeterminate	52/53/54/55	Toxic	Toxic
44	Lethal	Lethal	56	Mildly toxic	Lethal
51	Lethal	Lethal	57	Mildly toxic	No effect
50/51	Lethal	Lethal	58	Mildly toxic	No effect
44/50/51	Highly Lethal	No data	56/57/58	Lethal	Lethal
45	Toxic	Mildly Toxic	59/60	Toxic **	Lethal **

* Lethality conclusion based on failure to clone, when appropriate controls cloned readily; **Data not conclusive.

[46] **Genes 37-40:** Fragments carrying either of genes 37 or 39 could not be cloned in pPW19 in the expressed orientation, although each was cloned in the opposite orientation, suggesting that expression of each of these clones is lethal, even at the low level of expression without induction. Gene 39 was cloned in a more tightly controlled expression vector, and its expression was lethal.

[47] Expression of gene 40 by itself was only mildly toxic to either *E. coli* or *B. subtilis*. However, a plasmid expressing the triplet combination 38/39/40 could not be cloned in *E. coli*, except in one case where spontaneous integration of insertion sequence IS-1 had inactivated gene 40. In this plasmid, expression of the remaining wild-type genes 38/39 was lethal to *E. coli*. The same plasmid could not be transformed into *B. subtilis*, suggesting that even the uninduced level of 38/39 expression is

lethal to *B. subtilis*. The fact that gene 40 had to be inactivated to permit cloning in *E. coli* suggests that GP40 reinforces the lethal activity of GP38 and GP39.

[48] **Mutational analysis in Example 2 (FIG. 4)** shows that GP40 is essential for normal shutoff of host DNA synthesis, and that GP39 acts to restrain that activity of GP40. Mutational analysis (FIG. 3) also shows that both 39 and 40 are required for normal shutoff of host RNA and protein synthesis. The 38⁻ mutation was also deficient in shutoff of host DNA, RNA, and protein synthesis. Mutants inactivating gene 37 have also been isolated, but results with them are thus far inconclusive.

[49] **Gene 41:** Expression of gene 41 has only a mild effect on *E. coli*, but the plasmid carrying it could not be transformed into *B. subtilis*, suggesting that the low level of expression without induction is lethal to *B. subtilis*. The gene has apparently been cloned in *B. subtilis*, using a more tightly controlled vector, but experiments with that clone are thus far inconclusive. A mutation inactivating the gene is being isolated.

[50] **Genes 42 and 43:** Gene 42 showed no toxic effect on either bacterium, and results on gene 43 were too inconsistent to permit any conclusion.

[51] **Gene 44:** Expression of gene 44 was shown to be lethal to both *B. subtilis* and *E. coli*. FIG. 8 shows an example of killing of *E. coli* by expression of SPO1 gene 44. In cells growing in liquid medium, expression of the cloned gene 44 was induced at time 0. At subsequent times, the number of cells remaining viable was measured by plating in the absence of inducer. More than 99% of the cells were killed within three hours. Similar experiments showed the lethal effect of gene 39, gene 56 and two-gene operons 45/46 and 50/51.

[52] **Genes 44, 50, and 51:** Expression of either gene 44 or 51 alone is lethal to either *B. subtilis* or *E. coli*, inhibiting DNA, RNA, and protein synthesis. Adding gene 50 to the 51 clone increases its lethal efficiency, as does combining all 3 genes in a single clone. An *E. coli* mutant resistant to GP44 has a mutant *rpoB* gene, suggesting that the cellular target for GP44 is the bacterial RNA polymerase. Mutants resistant to genes 51/50 have been isolated, but the mutant gene(s) have not yet been identified. GP51 and GP44 each include a segment similar to a known RNA polymerase-binding motif, suggesting that GP51 may also act on RNA polymerase, although the RNA polymerase

(RNAP) mutation that protects against GP44 does not protect against GP51/50. The effects on DNA and protein synthesis are probably indirect effects of the inhibition of host RNA synthesis.

[53] We show that the products of SPO1 genes 44, 50, and 51 are required for the normal transition from early to middle gene expression during infection of *Bacillus subtilis* by bacteriophage SPO1; that they are also required for control of the shutoff of host DNA, RNA, and protein synthesis; and that their effects on host-shutoff could be accounted for by their effects on the regulation of gene expression. These three gene products had four distinguishable effects in regulating SPO1 gene expression: (1) GP44/50/51 acted to restrain expression of all SPO1 genes tested; (2) GP44 and/or GP50/51 caused additional specific repression of immediate-early genes; (3) GP44 and/or GP50/51 stimulated expression of middle genes; and (4) GP44 and/or GP50/51 stimulated expression of some delayed-early genes. Shutoff of immediate-early gene expression also required the activity of GP28, the middle gene-specific sigma factor. Shutoff of host RNA and protein synthesis was accelerated by either the 44- single mutant or the 50-51- double mutant, and more so by the 44-50-51- triple mutant. Shutoff of host DNA synthesis was accelerated by the mutants early in infection, but delayed by the 44-50-51- triple mutant at later times. Although GP50 is a very small protein, consisting almost entirely of an apparent membrane-spanning domain, it contributed significantly to each activity tested. The combination clone of genes 44/50/51 is the most efficiently lethal construct tested thus far (0.06% survival after 5 minutes induction).

[54] **Genes 45 and 46:** Expression of genes 45/46 is lethal to *B. subtilis* and highly toxic to *E. coli*. DNA, RNA, and protein synthesis are all affected about equally, showing a decrease of about 50% in the first hour, and no further decrease in the next 2 hours. No information is available yet on the cellular target, but resistant mutants should be readily obtainable.

[55] Individually, gene 46 is lethal but with diminished efficiency, and gene 45 is toxic but not lethal. Early results suggest that, during infection, a double mutant knocking out both genes is deficient in shutoff of host RNA and protein synthesis, but has no effect on host DNA synthesis.

[56] **Gene 47:** Gene 47 has little or no effect when expressed in either *B. subtilis* or *E. coli*.

[57] **Genes 48 and 49:** Gene 48 is strongly toxic but not lethal when expressed in either species. Although anomalies exist, the combination of genes 48 and 49 appears to be less toxic than 48 alone.

[58] **Genes 55-52:** Expression of gene 52 is lethal to *E. coli*. Wild-type gene 52 has not yet been tested on *B. subtilis*. Expression of a clone with genes 55/54/53 is toxic, but not lethal. In a single experiment, the latter inhibited bacterial protein synthesis without affecting RNA synthesis. Expression of the full operon is toxic, but not consistently lethal. This is apparently accounted for by a transcription terminator located between genes 53 and 52. Mutants resistant to GP52 have been isolated. The mutant gene(s) have not yet been identified, but while looking for them, several wild-type DNA segments whose over-expression protects against GP52 have been isolated. One of them includes a cluster of genes whose products are involved with protein synthesis, adding another suggestion that the gene 55 to 52 operon may target host protein synthesis. Gene 52 and gene 53 mutants are presently being isolated.

[59] **Genes 58-56:** Expression of the whole operon is lethal to both species. Expression of 56 alone is lethal to *B. subtilis*, specifically inhibiting cell division, without affecting DNA synthesis or chromosomal segregation. Infection by SPO1 inhibits cell division, and a gene 56 mutation prevents that inhibition. *B. subtilis* mutants resistant to GP56 have been isolated, but the mutant gene(s) has not yet been identified.

[60] **Genes 60-59:** The effects of GP59 and GP60 are uncertain because of apparent instability of the clone and inconsistency of data, but it appears that their expression may be lethal to *B. subtilis* in liquid culture but not on plates, while expression in *E. coli* is toxic, but not lethal.

[61] Since some of these genes can kill both gram-positive (*B. subtilis*) and gram-negative (*E. coli*) bacteria, they would probably be lethal to most bacterial pathogens. The first three of the lethal mechanisms studied targeted at least three different bacterial processes (RNA synthesis for gene 44, cell division for gene 56, and both DNA synthesis and RNA synthesis for genes 38, 39, and 40). The lethal mechanisms studied target a panoply of bacterial processes and are thus ideally suited for the development of antimicrobial agents.

EXAMPLE 2: MUTATIONAL ANALYSIS OF PROTEIN ROLE

[62] We have performed mutational analysis in order to determine what the roles of the 24 genes and their proteins are in the host takeover by SPO1.

[63] **Regulatory roles of genes 44, 50, and 51:** By site-specific mutagenesis, we have introduced a nonsense codon near the beginning of each of these genes, and have constructed all possible combinations of mutant genes. The nonsense mutations accelerated the rate of host-shutoff. FIG. 1 shows that shutoff of host RNA synthesis was accelerated by either the 44- single mutant or the 50-51- double mutant, and more so by the 44-50-51- triple mutant. Each data point in such an experiment represents the sum of host and phage RNA synthesis, so the decrease in total RNA synthesis by 12 minutes after infection provides a minimum estimate of the extent to which host RNA synthesis had been shut off. For wild-type and 44-50-51- these values were about 50% and 95%, respectively. The subsequent increases are due to increased phage RNA synthesis. Single mutants 50- and 51- had no significant effect (data not shown), showing that both GP50 and GP51 contribute to the activity of GP50/51.

[64] Each of the three mutations contributed to this effect. 44- and 50-51- each shut off host RNA synthesis more completely than wild-type, but less completely than the triple mutant. FIG. 7 shows that, while each of the 50- and 51- mutations had little effect by itself, each was necessary for the full effect of the triple mutant, 44-50-51-, and the double mutant, 50-51- (FIG. 7a and 7b)

[65] Similar results were seen when shutoff of host protein synthesis was measured by pulse-labeling with [³H]leucine, or by SDS-polyacrylamide gel electrophoresis of proteins pulse-labeled with [³⁵S]-labeled amino acids. Thus, GP44/50/51 restrained the normal shutoff of host gene expression, and each of genes 44, 50, and 51 contributed significantly to that activity.

[66] FIG. 2 shows that shutoff of host DNA synthesis was accelerated by the mutants early in infection, but delayed by the 44-50-51- triple mutant at later times. Only host DNA synthesis is assayed in such an experiment, since SPO1 incorporates hmUra into its DNA in place of thymine.

[67] Thus, GP44/50/51 delayed the shutoff of host RNA synthesis, and both GP44 and GP50/51 contributed significantly to that activity. GP44/50/51 delayed the early component of the shutoff of host DNA synthesis, while a later component required the activity of GP44 and/or GP50/51.

[68] **Independent Role of GP50:** GP50 includes only 23 amino acids, and consists almost entirely of an apparent membrane-spanning domain, with charged amino acids at each end. It is important to demonstrate that such an unusual protein actually plays a functional role. The first 8 nucleotides of gene 50 overlap the last 8 nucleotides of gene 51, and the mutation that inactivates gene 50 also destroys the termination codon of gene 51, permitting GP51 to run on for about 30 more amino acids, beyond its normal 131. Thus, demonstration of an independent role for GP50 depends on the effects of adding the gene 50 mutation to strains that already have gene 51 inactivated. This is shown in FIG. 7 and FIG. 12.

[69] **Regulation of SPO1 gene expression.** The negative effects of GP44/50/51 on shutoff of host RNA synthesis, and on the early phase of shutoff of host DNA synthesis, suggest that GP44/50/51 plays a regulatory role, rather than a directly causative role, in host-shutoff. We believe that we have identified the nature of that regulatory role, in that GP44/50/51 regulate the expression of all of the SPO1 genes known or expected to be involved in host-shutoff as shown in FIGs. 5 and 6. FIG. 5 shows that the triple mutant decreased transcription of at least some middle genes, while dramatically increasing transcription of all immediate-early genes. Thus the most obvious effect of GP44/50/51 is to facilitate the transition from immediate-early to middle gene expression.

[70] However, when the data from 44- and 50-51- are considered, it's clear that these three gene products cause at least 4 distinguishable effects, which may be seen in FIG. 6. The immediate-early genes shown are representative of all known immediate-early genes. All of the known delayed early genes are shown, and the middle genes shown were the only ones assayed. Gene 27 is expressed from both an early and a middle promoter.

[71] The first effect is the repression of all early and middle genes, which requires the activity of both GP44 and GP50/51. In comparison with wild-type, either the 44- mutation or the 50-51- double mutant caused increased expression of all genes tested, including immediate-early, delayed-

early, and middle genes. Therefore, GP44/50/51 caused repression of all of the SPO1 genes tested, and the full effect of this repression required the activity of both GP44 and GP50/51.

[72] The second effect is residual repression of immediate-early genes by either GP44 or GP50/51, even in the absence of the other. In comparison with 44- or 50-51-, the triple mutant 44-50-51- caused a further increase in expression of immediate-early genes. Therefore, there must be residual repressive activity present in the 44- or 50-51- strains, which is no longer there in 44-50-51-, and which therefore must be caused by GP50/51 in the 44- strain and by GP44 in the 50-51- strain.

[73] The third effect is stimulation of expression of middle genes by either GP44 or GP50/51. In comparison with 44- or 50-51-, the triple mutant caused a decrease in expression of middle genes. Therefore, there must be an activity present in the 44- or 50-51- strains, which stimulates middle gene expression. Since that activity is no longer there in 44-50-51-, it must be caused by GP50/51 in the 44- strain and by GP44 in the 50-51- strain.

[74] The fourth effect is stimulation of expression of certain delayed-early genes by either GP44 or GP50/51. In comparison with 44- or 50-51-, the triple mutant caused a decrease in expression of delayed-early genes 27 and 28. Therefore, there must be an activity present in the 44- or 50-51- strains, which stimulates delayed-early gene expression. Since that activity is no longer there in 44-50-51-, it must be caused by GP50/51 in the 44- strain and by GP44 in the 50-51- strain. With respect to delayed-early genes 37-40, such an activity can be seen for GP44 in the 50-51- strain, but not conclusively for GP50/51 in the 44- strain.

[75] These effects of GP44/50/51 on regulation of expression of early and middle genes provide a plausible explanation for the effects of GP44/50/51 on host shutoff. The effect on shutoff of host RNA synthesis can be explained if one or more of the immediate-early genes, regulated by GP44/50/51, specifies a product that causes shutoff of host RNA synthesis. The over expression of such genes, caused by the triple mutant, could cause the accelerated shutoff of host RNA synthesis that was observed in triple mutant infection. The intermediate levels of over expression, caused by mutants 44- and 50-51-, could account for their also causing intermediate levels of accelerated shutoff of host RNA synthesis. Similarly, the deficient shutoff of host DNA synthesis by the triple mutant at middle

times could be explained if that shutoff required one or more of the middle gene products whose expression is deficient in the triple mutant.

[76] Alternative explanations, in which GP44/50/51 have activities, in addition to their regulation of SPO1 gene expression, which directly affect host shutoff, remain possible, although intuitively less likely because of their additional complexity. A possible explanation of the second and third effects is that GP44 or GP50/51 facilitates the displacement of σ^A by GP28, but that is complicated by the fact that transcription of most host genes also requires σ^A . Under some conditions, rRNA synthesis is not shut off effectively by SPO1. Thus, a possible role of GP44/50/51 might be to prevent shutoff of rRNA synthesis, thereby accounting for some of the effect of 44-50-51- on RNA shutoff. All of these are potential subjects of future investigations.

[77] The simplest hypothesis as to how GP44/50/51 regulate gene expression is that they interact with the host's RNAP to change its transcriptional specificity. Our earlier results had already suggested that RNAP was the cellular target for these proteins. When gene 44 was expressed in uninfected cells of either *B. subtilis* or *E. coli*, it caused the shutoff of bacterial RNA synthesis and cell death (Wei and Stewart, 1993, 1995). DNA and protein synthesis were also shut off, presumably as indirect results of the effect on RNA synthesis. An *E. coli* mutation substituting valine for glutamate at position 1272 of the beta subunit of the RNAP provided resistance to the lethal effects of gene 44 (Sampath, unpublished results). When the gene 50/51 operon was expressed in uninfected cells, it also caused the shutoff of RNA, DNA, and protein synthesis and cell death (our unpublished results). The products of both genes 44 and 51 include acidic/hydrophobic domains similar to the domain in *E. coli* σ^{54} that is required for binding to RNAP, and one segment of GP51 shows substantial similarity to the product of SPO1 gene 27, which is required for normal transcription of late genes (Greene *et al.*, 1982; Stewart, 1993). Because of the very small size of GP50 (23 amino acids), its presence in the same operon with, and overlapping gene 51, and the synergistic effects of mutations in genes 50 and 51 we hypothesize that GP50 participates in the same activities as GP51. We suppose that the observed effects of GP44 or GP50/51 in uninfected cells are incidental consequences of their binding to RNAP, which causes effects in the opposite direction when in the context of phage infection.

[78] **Causative roles of genes 37, 38, 39, and 40 in host-shutoff.** Nonsense mutations were introduced into each of genes 37 to 40 as described above. FIG. 3 shows that both the 40- and the 39- mutants are deficient in shutoff of host RNA synthesis, that 40- has the greater effect, and that the effects of the two mutations are not additive. These mutants had similar effects on shutoff of host protein synthesis (data not shown). However, as shown in FIG. 4, 40- inhibited but 39- accelerated the shutoff of host DNA synthesis, and 40- suppressed this effect of 39-. Apparently GP39 restrains the effect of GP40 on shutoff of host DNA synthesis, while both work in the same direction on host RNA synthesis. Both shutoffs occur so rapidly that it's unlikely that either effect of the 40- mutation is an indirect effect of the other. Genes 37-40 comprise one operon, with 38 and 39 overlapping slightly, suggesting the possibility of translational coupling and joint activity (Stewart *et al.*, 1998). Preliminary results are consistent with GP37 and GP38 also participating in the same shutoffs. Thus, GP40 is required for normal shutoff of host DNA and RNA synthesis, and each of the other genes in this operon does or may play a significant role in each of those shutoffs.

[79] **Onset of Phage DNA Synthesis:** FIG. 9 shows the effect of various of the mutant combinations on the onset of phage DNA synthesis. Each mutant combination that caused substantial delay in shutoff of host DNA synthesis also caused delay in phage DNA synthesis. This is probably because the continued use of cellular resources for host DNA synthesis meant that they were not fully available for phage DNA synthesis. Also contributing in most cases would be the diminished activity of phage genes required for phage DNA replication.

[80] **Single Step Growth Curves:** FIG. 10 and 11 show the effects of the various mutations on single step growth curves. Despite the profound effect of several of the multiple mutants on phage gene expression and DNA replication, production of progeny phage was only moderately affected. Each mutant that caused significant delay in phage DNA synthesis also caused a significant extension of the latent period. Only those mutants with a deficiency in phage DNA synthesis caused decreases in burst size. These were less than proportional to the decreases in DNA synthesis, suggesting that phage DNA is produced in excess in wild-type infections under these conditions.

EXAMPLE 3: CONTINUED ANALYSIS

[81] For those genes whose lethality is not yet certain, we will determine the effects of gene expression unequivocally by cloning in more tightly controlled expression vectors. For each of these lethal genes and gene combinations, we will attempt to determine the mechanism by which it exerts its lethal effect.

[82] To determine the cellular targets of the lethal proteins, we will select bacterial mutants resistant to each of the lethal genes, by plating, under inducing conditions, the strain carrying the lethal gene, and identifying those colonies whose resistance is caused by a chromosomal mutation. Each mutant gene will be identified, by screening a library of the chromosomal DNA for capacity to protect against the lethal effect of the phage gene. The methodology used will be similar to that described in Wei and Stewart (1995). Mutants resistant to 4 of the lethal genes have already been isolated, and in this manner the gene 44- resistant mutation was determined to be in the *rpoB* gene, specifying the B subunit of the RNA polymerase.

[83] In case the protective mutation cannot be located by such a library screen (for instance because the protective mutation proves to be recessive, or because the protective gene proves to be unclonable), we will locate the protective mutation by genetic mapping. For *E. coli*, we will determine the approximate location by co-transfer with the Tn10 markers of the set of Hfr strains developed by Wanner (1986) and available from the *E. coli* Genetic Stock Center, and then determine the location more precisely by degree of linkage in P1 transduction (procedures as in Miller, 1992). For *B. subtilis*, we will determine the approximate location by showing linkage in PBS1 transduction with markers represented in the Dedonder *et al.* (1977) or Zahler (Vandeyar and Zahler, 1986) mapping kits, available from the *Bacillus* Genetic Stock Center. For finer structure mapping, we will look for the degree of co-transformation with markers within the region that showed high frequency co-transduction. The procedures are modifications of those described in Harwood and Cutting (1990). The availability of a complete set of mutants inactivating each gene in the *B. subtilis* genome (Kobayashi *et al.*, 2003) will permit precise location of each affected gene, which will then be confirmed by nucleotide sequencing.

[84] **Identification of protein:protein interactions:** To confirm the interaction of the lethal proteins with the targets thus tentatively identified, and to identify targets not found by such

mutational analysis, host proteins that interact with each lethal SPO1 protein will be identified. We will make use of several genetic and physical techniques for identifying protein-protein interactions. Since such techniques can produce both false positives and false negatives (Legrain, 2002), we will use two or more of these techniques with each of the proteins. These techniques include:

[85] **Identification of cellular targets by genetic selection:** Since these SPO1 proteins interfere with essential host processes, many of them are lethal when expressed individually in uninfected bacteria. Thus, their cellular targets may be identified by selecting bacterial mutants resistant to their lethal effects. Most of the proteins have similar effects on *E. coli* and *B. subtilis* (which share substantial homology), permitting the use of the more versatile *E. coli* cloning systems for initial identification of certain cellular targets. Targets initially identified in *E. coli* will be confirmed by testing the effect of the *B. subtilis* homolog.

[86] *E. coli* mutants resistant to the lethal effect of GP44 were selected by plating under inducing conditions. One mutated gene identified was the *rpoB* gene (which specifies the beta subunit of RNAP), by screening a genomic library for capacity to protect against the lethal effect (Wei and Stewart, 1995; Sampath, unpublished results). Although alternative explanations are possible, this strongly suggests that GP44 targets the host RNAP. Similar analyses are in progress with others of the SPO1 lethal genes.

[87] **Pull-down assays.** Proteins to be tested will either be expressed as glutathione-S-transferase (GST) fusions for attachment to glutathione sepharose beads (Pharmacia) or directly attached to N-hydroxysuccinimide (NHS)-activated sepharose resin (Pharmacia). The coupling of the NHS to primary amine is done under non-denaturing conditions at room temperature, and is quite efficient. Extracts of either infected or uninfected *B. subtilis* or *E. coli* cultures will be passed over the column, and the retained proteins will be eluted using increasing salt concentrations or urea denaturation. The eluant will then be subjected to SDS-PAGE, and the proteins electroeluted onto PVDF membranes for N-terminal sequencing and identification (Einarson and Orlinick, 2002).

[88] **Phage Display.** Because only a few amino acids from one protein are needed to provide effective binding to another protein (Scott, 2001), it is not necessary to probe a library of whole proteins to identify binding partners. A phage display library (Willats, 2002; Barbas *et al.*, 2001)

displaying random sequence peptides 5 to 7 amino acids long will include most of the sequences necessary to confer specificity of binding to any protein. When a single protein is used to probe such a library, it can potentially identify the binding sequences of most or all proteins capable of binding to the probe protein. For instance, Malys *et al.* (2002) probed a random sequence peptide library with the T4 terminase protein, and identified all proteins previously known to bind to that protein, plus five other proteins whose binding was not previously known, but was subsequently confirmed in the one case conclusively tested.

[89] Each of the proteins will be used to probe a random sequence peptide library, displayed on either phage T4 or phage M13. Several M13 libraries are available from New England Biolabs and have been used successfully in a long list of experiments (NEB, 2003). The peptide sequences identified as binding to each of the probe proteins will be used to probe the database of all *B. subtilis* and SPO1 gene products, thus identifying those proteins having segments capable of binding to the probe protein. The *B. subtilis* and *E. coli* genomes are completely known, and the virtually complete SPO1 sequence is also available (Hendrix *et al.*, 2003).

[90] **Immunoprecipitation.** Antibodies will be prepared against each of the proteins. Each antibody will be used to precipitate the targeted protein from extracts of infected cells, following procedures as described (Wei and Stewart, 1993; Wei and Stewart, 1995). The proteins in the precipitate will be identified using the same procedure as described.

[91] **Two-hybrid analysis.** We anticipate that the above procedures will be sufficient to identify all or most of the proteins that interact with any of the lethal SPO1 proteins. However, if further information is needed, we will make use of a two-hybrid system. Noirot-Gros *et al.* (2002) described the use of yeast two-hybrid screens to identify many interactions between *B. subtilis* proteins, and this procedure can readily be adapted to interactions between SPO1 proteins and bacterial proteins.

EXAMPLE 4: NONLYTIC PHAGE THERAPY

[92] Certain pathogens (e.g. *E. coli* O157:H7 or *Neisseria meningitidis*) cannot be treated with lytic phage or conventional antibiotics, because rapid destruction of the bacteria causes the precipitate release of lethal toxins that can kill the patient. The antimicrobial genes and peptides described herein solve this problem because each of them kills *without* lysing the bacterial cell. This has

been documented directly for several of the genes by showing either microscopically or turbidimetrically that the structure of the cells remains intact even after they have lost viability. Maintenance of structural integrity can also be inferred for almost all of them, since all but gene 52 are expressed early during infection and expression of all early genes is insufficient to cause lysis during infection. Thus, the proteins could be used to treat such infections directly, or a temperate bacteriophage could be used to target the genes to the bacteria.

[93] A transducing bacteriophage, specific for the particular pathogen, would introduce the lethal SPO1 gene(s), either on a plasmid that replicates in that bacterium or as part of a structure that would integrate into the bacterial genome. Exponential Biotherapies Inc. has successfully completed Phase 1 clinical trials of one of their first-generation phage therapy vehicles, a lytic phage that destroys pathogenic *Enterococcus* bacteria. They have patents on a method that selects for phages that survive in the circulation long enough to rescue animals from otherwise-lethal bacteremias (US20010026795 and US20010043924). The antimicrobial SPO1 genes would be used in the second generation of phage therapy vehicles, in which individual lethal genes or gene combinations would be delivered to the infecting bacteria by a transducing phage.

EXAMPLE 5: PEPTIDOMIMETICS

[94] To be effective therapeutically, the antibiotic activities characteristic of these various lethal proteins must be delivered to the infecting bacterial pathogen. This could be accomplished in any of several ways, one of which is to design small molecule drugs that mimic the active sites of the proteins. To be effective, the antibiotic characteristic of various lethal proteins must be delivered to the infecting bacterial pathogen, in concentrations high enough to be therapeutic. By designing small molecule drugs that mimic the active sites of the lethal proteins, but that have favorable pharmacological characteristics with regard to delivery, uptake, human toxicity, and potency, a directed peptidomimetic molecule can be designed. Examples of successful use of such peptidomimetic small molecules include: Tian *et al.*, 1998; Salvemini *et al.*, 1999, 2002; Andrade-Gordon *et al.*, , 1999; Wrighton and Gearing, 1999; Zhang, 2000; Ohkanda *et al.*, 2001; Smith *et al.*, 2001; Air *et al.*, 2002; Gadek *et al.*, 2002; Fotsch *et al.*, 2003; and Martin *et al.*, 2003.

[95] Structures of the lethal proteins will be determined. The lethal SPO1 proteins are relatively small (ranging from 77 to 255 amino acids) so we can determine their structures by X-ray crystallography and/or NMR spectroscopy. Each functional site will be identified by 1) mutagenesis of the cloned gene, 2) selection for mutants that have lost their lethal activity, and 3) identification of the positions of the mutant amino acids in the 3-dimensional structures. Similarly, the structures of the target molecules will be determined, and the targeted surfaces identified by the positions of mutations causing resistance to the lethal proteins. Where possible, we will co-crystallize the lethal protein with the target molecule, to determine changes in structure caused by their interaction.

[96] Small molecules can then be designed that attempt to mimic the structure of the active site. High throughput screening of many candidate molecules for binding to the target protein or inhibition of its function, depending on the specific target identified, will provide lead molecules for peptidomimetic chemicals.

[97] **Preparation for X-ray crystallography.** The lethal SPO1 proteins are relatively small (ranging from 77 to 255 amino acids) and their origins in a prokaryotic host make it likely that the proteins will be correctly folded on expression. Each of the purified proteins will be tested by light scattering and native gel electrophoresis to determine whether the samples are monodisperse. Most proteins will be tested for proper folding, by taking their Heteronuclear-Single Quantum Correlation (HSQC) spectra, using either the 500 or 600 MHz NMR at Rice University. To do this, the proteins will be ^{15}N -labeled by growing cells in minimal media supplemented with ^{15}N -ammonium chloride. Proteins that are partially unfolded can be rapidly identified by ^{15}N -HSQC NMR spectra, and the salt, pH and ligand conditions can be adjusted to provide a homogeneous, well-folded sample suitable for X-ray diffraction studies. Proteins that are well-folded and active will then be subjected to a sparse-matrix crystallization protocol to identify conditions appropriate for macromolecular crystal growth. The smaller proteins such as GP46 and GP56 would also be amenable to structure determination by NMR. They will be isotopically labeled with ^{15}N or ^{15}N and ^{13}C and their structures determined by multidimensional heteronuclear NMR methods. In cases where a toxic protein is expressed in limited quantities, we will skip the NMR screening and proceed directly to crystallography.

[98] **High-throughput crystallographic screening.** The Rice University Crystallography Core Facility uses a MATRIX TECHNOLOGIES® HYDRA II-PLUS ONE™ liquid-handling robot and automated imaging microscope, for high-throughput crystallographic screening (Diversified Scientific). Such facilities give us the ability to rapidly screen far more crystallographic conditions with far less material than traditional vapor diffusion methods. This is particularly useful for proteins whose yield has been limited due to their toxic nature. Using our high-throughput facilities, 1 mg of protein at 10 mg/ml can be subjected to 500 trial conditions, as compared to 100 using manual methods. Analysis is facilitated by construction of relational databases for correlation of high-throughput crystallization data with incomplete-factorial sparse matrix generation and development of new second generation matrices derived from the observed trials. Recent studies from the NIH structural genomics initiative have suggested that in 90% of cases of a monodisperse and well-folded protein where a crystallization condition is found, the condition was found in only ~200-500 trials. Failure to obtain crystals with good diffraction properties after 500 trials suggests that either addition of a ligand/cofactor or partial proteolysis studies may be appropriate for further trials. Once crystallization conditions have been identified, structures will be determined using either Multi-wavelength Anomalous Diffraction (MAD) (Hedrickson and Ogata, 1997) or Multiple Isomorphous Replacement (MIR) (McPherson, 1999). Model building and refinement (Shamoo *et al.*, 1995, Shamoo and Steitz, 1999) provides additional information from small sample sizes and allows generation of ideal peptidomimetic structures.

[99] **Mutagenesis.** To map the functional site of each of the lethal proteins, genes encoding the lethal proteins will be subjected to error prone PCR *in vitro* (Spee *et al.*, 1993; Miyazaki *et al.*, 2000) and then reintroduced back into either *E. coli* or *B. subtilis* by transformation. Mutations that strongly reduce or abolish host lethality will be selected by survival of the host to form colonies. Sequencing of the mutants will permit mapping the changes onto the protein's primary structure, and thus onto the three-dimensional structure as determined above, hopefully providing a tentative identification of the active site.

[100] **Structure of target molecules and target sites.** It should be possible to identify the target site within the target molecule, by identifying the amino acids whose mutation protects the target against the activity of the lethal protein, and finding which of those amino acids in the wild-type protein form a surface complementary to the binding surface of the lethal protein. Thus, when a target

has been identified, the wild-type gene specifying it will be cloned and subjected to mutagenesis by error prone PCR. The library consisting of that mutagenized population will be transformed back into the host cells, selecting for mutants resistant to the lethal protein. The position of the mutations will be determined by nucleotide sequencing, and those positions mapped onto the 3-dimensional structure. In cases where the target structure is not known, we will determine it whenever possible. Such analysis will allow us to determine a possible mechanism of action; for example resistant mutations may map to a known protein-protein contact surface or active site. We can then develop an accurate structure-function characterization of the host-SPO1 protein interaction and mechanism.

[101] To minimize the likelihood of toxic side effects when the peptidomimetic molecules are introduced into clinical use, our highest priorities for the above analyses will be those of the lethal SPO1 proteins that do not target a protein with an obvious human homolog. However, even those whose target does have a human homolog might be worth pursuing, since the target may function differently or be less accessible in human cells.

EXAMPLE 6: ANTIMICROBIAL TREATMENTS

[102] Information already in hand about some of these lethal proteins shows that they target a variety of different bacterial processes, so we expect to develop a panoply of different antibiotic mechanisms, each targeting a different bacterial process. Since none of the lethal gene products show substantial homology to any known protein, it is highly probable that each mechanism will be novel, meaning that bacteria resistant to currently used antibiotics should be susceptible to each of the antibiotics based on these lethal proteins.

[103] The antibiotics to be developed would have the following advantages: (1) targeting of a wide spectrum of gram-positive and gram-negative bacterial pathogens; (2) effectiveness against anthrax, plague, and other anticipated agents of biological warfare; (3) effectiveness against bacteria that are resistant to presently known antibiotics; (4) inclusion of a diverse spectrum of antibiotic mechanisms; and (5) minimal release of lethal toxins during treatment.

[104] Combining two or more lethal genes in the same clone can form more efficiently lethal constructs. For instance, the triplet combination of genes 44, 50, and 51 kills more than 20 times

as efficiently as either of its components cloned individually in the same vector (0.06% survival after 5 minutes induction).

[105] Since few, if any, existing antibiotics are based on a specific bacteriophage gene, and since few, if any, of the lethal SPO1 gene products show substantial homology to any protein presently in the public database, it is likely that each lethal mechanism will be novel. This should help avoid the resistance mechanisms of the current generation of resistant bacteria.

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